Amendments to the Specification

In the Brief Description of the Drawings, please make the following changes:

Replace the paragraph beginning at page 22, line 11, with the following paragraph:

FIGURE 1A-C. Forms of Immune Augmentation. This figure demonstrates an overview of various embodiments of the invention, including "immunoadjuvant" embodiments wherein the immunopotentiating pootein is simply admixed with a compound against which an immune response is desired (top panel), or where and an actual heteroconjugate is formed between the immunopotentiating protein and the compound (middle panel). In the bottom panel is shown an embodiment wherein the immunopotentiating ligand is actually a bifunctional cojugate conjugate formed between two antibodies.

Replace the paragraph beginning at page 22, line 22, with the following paragraph:

FIGURE 2<u>A-1-B-4</u>. Activation of peripheral lymph node T cells from anti-CD3-treated C3H mice as assessed by flow cytometry (FCM). Two color FCM from control animals and those treated with, 4, 40, or 400 μg of anti-CD3 are displayed as contour plots on a logarithmic scale. Intensity of green FITC fluorescence is plotted along the x-axis and red (B-phycoerythrin) fluorescence is plotted along the 7-axis. (A) Anti-CD4 staining on the x-axis and anti-IL-2 receptor (I1-2R) staining on the y-axis. (B) Anti-CD3 staining on the x-axis and anti-Thy-1 staining on the y-axis. C3H/HeN MTV mice were killed 18 hours after intravenous injection of purified anti-CD3 (MAb 145-2C11) that was grown and purified as described (24). Femoral, axillary, and mesenteric lymph nodes were removed and dissociated into a single-cell suspension and FCM analysis was performed (25). Cells were stained with FITC-anti-CD3 or FITC-anti-CD4 (MAb GK1.5) (Becton Dickinson), and biotin-conjugated anti-IL-2R (MAb 3C7) or biotin-

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conjugated MAb to Thy-1.2 (Becton Dickinson), then counterstained with B-phycoerythrin-conjugated egg white avidin (Jackson Immuno Research Laboratories). These results show that low dose (4 ug) anti-CD3 treatment activates T cells as evidenced by IL-2R expression but does not modulate T cell receptors.

Replace the paragraph beginning at page 23, line 33, with the following paragraph:

FIGURE 4A-B. Colony Stimulating Factor (CSF) in serum of mice after injection of anti-CD3. Pooled sera from three animals were placed at 6% final dilution with murine bone marrow cells, and the number of colonies were counted after 7 days. Each sample was tested in duplicate and the results were averaged. In all cases, duplicate values differed by no more than 5%. A: Mice received 400µg anti-CD3 Ig(•), or 250µg of F(ab')₂ fragments of anti-CD3 (o). (The number of colonies at 3h for anti-CD3 treated mice was more than 300.) B: Number of colonies after various doses of anti-CD3. Serum was collected 3 h after injection. These results show that anti-CD3 in vivo induces lymphokines including colony stimulating factors.

Replace the paragraph beginning at page 24, line 12, with the following paragraph:

FIGURE 5A-B. Clinical Response to anti-CD3 (OKT3) Treatment. A: Increased allogenic MHC response in patients treated with OKT3. B: Proliferation of T cells before and after OKT3 treatment in the presence (cross-hatched bars) or absence (closed bars) of rIL-2 suggest that in vivo treatment with OKT3 activates human T cells.

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Replace the paragraph beginning at page 24, line 34, with the following paragraph:

FIGURE 87 <u>8A-B</u>. IL-2R expression on T cells from SEB-treated mice. Mice were treated with increasing doses of SEB (0, 5, 50, 250µg). I1-2R expression after 18 hours was compared using flow cytometry and showed enhanced expression. Dose response was observed.

Replace the paragraph beginning at page 25, line 12, with the following paragraph:

FIGURE 10A-1-B-3. Expansion of $V_{\beta}8^{+}$ cells in SEB-treated mice. Three days after treatment of mice with SEB, spleen cells were incubated with anti- $V_{\beta}8$ and $V_{\beta}8^{+}$ cells were assayed by flow cytometry. Expansion of $V_{\beta}8^{+}$ cells was observed due to SEB treatment.

Replace the paragraph beginning at page 27, line 13, with the following paragraph:

Panel A: IgG anti-FITC antibody production in anti-CD3 treated mice immunized with FITC-BSA in complete Freunds adjuvant (CFA) or PBS, compared to ELISA measurements of sera from control mice (open bars). Panel B: IgG anti-FITC antibody production measured from sera of FITC-anti-CD3 treated mice (left-hatched bars) compared to FITC-normal Hamster Ig (cross-hatched bars) measured by ELISA performed on day 10 bleeds.

Please replace the Abstract with the amended Abstract attached hereto.